

K955651 MEDTRONIC SHERPA GUIDING CATHETERJan 22, 1996
41 days to decisionK955651 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k955651/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Dec 12, 1995
Decision date	Jan 22, 1996
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronics Interventional Vascular
Location	Danvers, MA, US
Contact	JOSEPH O MAGLIOZZI
510(k) history	21 submissions · 21 cleared · 1992-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955651/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026